

Chia-Ron Yang

List of Publications by Year in descending order

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Version: 2024-02-01

41
papers

1,412
citations

361388

20
h-index

330122

37
g-index

43
all docs

43
docs citations

43
times ranked

2053
citing authors

#	ARTICLE	IF	CITATIONS
1	High glucose-induced apoptosis in human vascular endothelial cells is mediated through NF- κ B and c-Jun NH2-terminal kinase pathway and prevented by PI3K/Akt/eNOS pathway. Cellular Signalling, 2006, 18, 391-399.	3.6	223
2	Soluble Decoy Receptor 3 Induces Angiogenesis by Neutralization of TL1A, a Cytokine Belonging to Tumor Necrosis Factor Superfamily and Exhibiting Angiostatic Action. Cancer Research, 2004, 64, 1122-1129.	0.9	107
3	Decoy Receptor 3 Increases Monocyte Adhesion to Endothelial Cells via NF- κ B-Dependent Up-Regulation of Intercellular Adhesion Molecule-1, VCAM-1, and IL-8 Expression. Journal of Immunology, 2005, 174, 1647-1656.	0.8	91
4	Histone deacetylase inhibitors increase microRNA-146a expression and enhance negative regulation of interleukin-1 β signaling in osteoarthritis fibroblast-like synoviocytes. Osteoarthritis and Cartilage, 2013, 21, 1987-1996.	1.3	87
5	(<i>N</i> -Hydroxycarbonylbenzylamino)quinolines as Selective Histone Deacetylase 6 Inhibitors Suppress Growth of Multiple Myeloma <i>in Vitro</i> and <i>in Vivo</i> . Journal of Medicinal Chemistry, 2018, 61, 905-917.	6.4	69
6	The novel histone de acetylase 6 inhibitor, MPTOG211, ameliorates tau phosphorylation and cognitive deficits in an Alzheimer's disease model. Cell Death and Disease, 2018, 9, 655.	6.3	68
7	Aciculatin inhibits lipopolysaccharide-mediated inducible nitric oxide synthase and cyclooxygenase-2 expression via suppressing NF- κ B and JNK/p38 MAPK activation pathways. Journal of Biomedical Science, 2011, 18, 28.	7.0	62
8	5-Aroylindoles Act as Selective Histone Deacetylase 6 Inhibitors Ameliorating Alzheimer's Disease Phenotypes. Journal of Medicinal Chemistry, 2018, 61, 7087-7102.	6.4	56
9	Intracerebral transplantation of neural stem cells combined with trehalose ingestion alleviates pathology in a mouse model of Huntington's disease. Journal of Neuroscience Research, 2009, 87, 26-33.	2.9	49
10	Denbinobin upregulates miR-146a expression and attenuates IL-1 β -induced upregulation of ICAM-1 and VCAM-1 expressions in osteoarthritis fibroblast-like synoviocytes. Journal of Molecular Medicine, 2014, 92, 1147-1158.	3.9	46
11	Indole-3-ethylsulfamoylphenylacrylamides: Potent histone deacetylase inhibitors with anti-inflammatory activity. European Journal of Medicinal Chemistry, 2014, 85, 468-479.	5.5	41
12	Decoy Receptor 3 Expression in AsPC-1 Human Pancreatic Adenocarcinoma Cells via the Phosphatidylinositol 3-Kinase-, Akt-, and NF- κ B-Dependent Pathway. Journal of Immunology, 2008, 181, 8441-8449.	0.8	34
13	3-Aroylindoles display antitumor activity <i>in vitro</i> and <i>in vivo</i> : Effects of N1-substituents on biological activity. European Journal of Medicinal Chemistry, 2017, 125, 1268-1278.	5.5	34
14	Preclinical anti-arthritic study and pharmacokinetic properties of a potent histone deacetylase inhibitor MPTOG009. Cell Death and Disease, 2014, 5, e1166-e1166.	6.3	31
15	Anti-metastatic activity of MPTOG211, a novel HDAC6 inhibitor, in human breast cancer cells <i>in vitro</i> and <i>in vivo</i> . Biochimica Et Biophysica Acta - Molecular Cell Research, 2019, 1866, 992-1003.	4.1	31
16	Anti-Arthritic Effects of Magnolol in Human Interleukin 1 β -Stimulated Fibroblast-Like Synoviocytes and in a Rat Arthritis Model. PLoS ONE, 2012, 7, e31368.	2.5	28
17	2-(Phenylsulfonyl)quinoline N-hydroxyacrylamides as potent anticancer agents inhibiting histone deacetylase. European Journal of Medicinal Chemistry, 2016, 122, 92-101.	5.5	28
18	Anticancer activity of MPTOG157, a derivative of indolylbenzenesulfonamide, inhibits tumor growth and angiogenesis. Oncotarget, 2015, 6, 18590-18601.	1.8	26

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19	Neuroprotective Studies of Evodiamine in an Okadaic Acid-Induced Neurotoxicity. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5347.	4.1	26
20	Potent Anti-Inflammatory Effects of Denbinobin Mediated by Dual Inhibition of Expression of Inducible No Synthase and Cyclooxygenase 2. <i>Shock</i> , 2011, 35, 191-197.	2.1	25
21	Denbinobin suppresses breast cancer metastasis through the inhibition of Src-mediated signaling pathways. <i>Journal of Nutritional Biochemistry</i> , 2011, 22, 732-740.	4.2	21
22	1- <i>N</i> -hydroxyacrylamide)indolines Histone Deacetylase Inhibitors Are Potent Cytokine Release Suppressors. <i>ChemBioChem</i> , 2013, 14, 1248-1254.	2.6	21
23	The anticancer effects of MPTOG211, a novel HDAC6 inhibitor, combined with chemotherapeutic agents in human acute leukemia cells. <i>Clinical Epigenetics</i> , 2018, 10, 162.	4.1	20
24	THIAZOLIDINEDIONES INHIBIT TNF- α -MEDIATED OSTEOCLAST DIFFERENTIATION OF RAW264.7 MACROPHAGES AND MOUSE BONE MARROW CELLS THROUGH DOWNREGULATION OF NFATc1. <i>Shock</i> , 2010, 33, 662-667.	2.1	19
25	Novel histone deacetylase inhibitor MPTOG009 induces cell apoptosis and synergistic anticancer activity with tumor necrosis factor-related apoptosis-inducing ligand against human hepatocellular carcinoma. <i>Oncotarget</i> , 2016, 7, 402-417.	1.8	19
26	Moscatalin Ameliorates Tau Phosphorylation and Cognitive Deficits in Alzheimer's Disease Models. <i>Journal of Natural Products</i> , 2019, 82, 1979-1988.	3.0	18
27	MPTOG413, A Novel HDAC6-Selective Inhibitor, and Bortezomib Synergistically Exert Anti-tumor Activity in Multiple Myeloma Cells. <i>Frontiers in Oncology</i> , 2019, 9, 249.	2.8	18
28	A novel dual HDAC and HSP90 inhibitor, MPTOG449, downregulates oncogenic pathways in human acute leukemia in vitro and in vivo. <i>Oncogenesis</i> , 2021, 10, 39.	4.9	15
29	Combined treatment with Denbinobin and Fas ligand has a synergistic cytotoxic effect in human pancreatic adenocarcinoma BxPC-3 cells. <i>British Journal of Pharmacology</i> , 2009, 157, 1175-1185.	5.4	14
30	Inhibition of neuronal migration by JONES antibody is independent of 9-O-acetyl GD3 in GD3-synthase knockout mice. <i>Journal of Neuroscience Research</i> , 2007, 85, 1381-1390.	2.9	13
31	Anti-Inflammatory and Tau Phosphorylation Inhibitory Effects of Eupatin. <i>Molecules</i> , 2020, 25, 5652.	3.8	13
32	Identification of a dual TAOK1 and MAP4K5 inhibitor using a structure-based virtual screening approach. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021, 36, 98-108.	5.2	10
33	Aciculin Inhibits Granulocyte Colony-Stimulating Factor Production by Human Interleukin 1 β -Stimulated Fibroblast-Like Synoviocytes. <i>PLoS ONE</i> , 2012, 7, e42389.	2.5	9
34	Identification and analysis of a selective DYRK1A inhibitor. <i>Biomedicine and Pharmacotherapy</i> , 2022, 146, 112580.	5.6	8
35	4-Substituted 2-amino-3,4-dihydroquinazolines with a 3-hairpin turn side chain as novel inhibitors of BACE-1. <i>Bioorganic Chemistry</i> , 2020, 95, 103135.	4.1	7
36	An oral quinoline derivative, MPTOB392, causes leukemic cells mitotic arrest and overcomes drug resistant cancer cells. <i>Oncotarget</i> , 2017, 8, 27772-27785.	1.8	6

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37	Discovery of a novel cyclin-dependent kinase 8 inhibitor with an oxindole core for anti-inflammatory treatment. <i>Biomedicine and Pharmacotherapy</i> , 2022, 146, 112459.	5.6	5
38	O-methylated flavonol as a multi-kinase inhibitor of leukemogenic kinases exhibits a potential treatment for acute myeloid leukemia. <i>Phytomedicine</i> , 2022, 100, 154061.	5.3	5
39	Anti-leukemia effects of the novel synthetic 1-benzylindole derivative 21-900 in vitro and in vivo. <i>Scientific Reports</i> , 2017, 7, 42291.	3.3	4
40	Structure-based virtual screening and biological evaluation of novel small-molecule BTK inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022, 37, 226-235.	5.2	3
41	Anti-metastatic evaluation of a novel HDAC6 inhibitor YH308 in human breast cancer cells in vitro and in vivo. <i>Proceedings for Annual Meeting of the Japanese Pharmacological Society</i> , 2018, WCP2018, PO4-6-38.	0.0	0